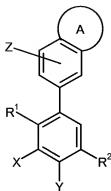


**Amendments to the claims**

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended) A compound of formula (I):



(I)

wherein

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur, optionally substituted by up to two substituents independently selected from C<sub>1-6</sub>alkyl, -(CH<sub>2</sub>)<sub>k</sub>-C<sub>3-7</sub>cycloalkyl, halogen, -CN, trifluoromethyl, -(CH<sub>2</sub>)<sub>k</sub>OR<sup>3</sup>, -(CH<sub>2</sub>)<sub>k</sub>CO<sub>2</sub>R<sup>3</sup>, -(CH<sub>2</sub>)<sub>k</sub>NR<sup>3</sup>R<sup>4</sup>, -(CH<sub>2</sub>)<sub>k</sub>CONR<sup>3</sup>R<sup>4</sup>, -(CH<sub>2</sub>)<sub>k</sub>NHCOR<sup>3</sup>, -(CH<sub>2</sub>)<sub>k</sub>SO<sub>2</sub>NR<sup>3</sup>R<sup>4</sup>, -(CH<sub>2</sub>)<sub>k</sub>NHSO<sub>2</sub>R<sup>3</sup>, -(CH<sub>2</sub>)<sub>k</sub>SO<sub>2</sub>(CH<sub>2</sub>)<sub>m</sub>R<sup>5</sup> [or] a 5- or 6-membered heterocyclyl ring containing nitrogen optionally substituted by C<sub>1-2</sub>alkyl or -(CH<sub>2</sub>)<sub>k</sub>CO<sub>2</sub>R<sup>3</sup>, and a 5-membered heteroaryl ring optionally substituted by C<sub>1-2</sub>alkyl; or

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur substituted by [[-BR<sup>6</sup>]] -B<sup>1</sup>R<sup>6</sup>, and A is optionally further substituted by one substituent selected from -OR<sup>7</sup>, halogen, trifluoromethyl, -CN, -CO<sub>2</sub>R<sup>7</sup> and C<sub>1-6</sub>alkyl optionally substituted by hydroxy; or

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur substituted by -(CH<sub>2</sub>)<sub>n</sub>heterocyclyl wherein the heterocyclyl is a 5- or 6-membered heterocyclic ring containing one or two heteroatoms independently selected from oxygen, sulfur and nitrogen optionally substituted by up to two substituents independently selected from oxo, C<sub>1-6</sub>alkyl, OR<sup>7</sup>, -NR<sup>7</sup>R<sup>8</sup> and -CONR<sup>7</sup>R<sup>8</sup>, and A is optionally further substituted by one substituent selected from -OR<sup>7</sup>, halogen, trifluoromethyl, -CN, -CO<sub>2</sub>R<sup>7</sup> and C<sub>1-6</sub>alkyl optionally substituted by hydroxy; or

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur substituted by -(CH<sub>2</sub>)<sub>q</sub>aryl or

-(CH<sub>2</sub>)<sub>q</sub>heteroaryl wherein the aryl or heteroaryl is optionally substituted by one or more substituents independently selected from oxo, C<sub>1-6</sub>alkyl, halogen, -CN, trifluoromethyl, -OR<sup>9</sup>, -(CH<sub>2</sub>)<sub>t</sub>CO<sub>2</sub>R<sup>10</sup>, -NR<sup>9</sup>R<sup>10</sup>, -(CH<sub>2</sub>)<sub>t</sub>CONR<sup>9</sup>R<sup>10</sup>, -NHCOR<sup>9</sup>, -SO<sub>2</sub>NR<sup>9</sup>R<sup>10</sup>, -NHSO<sub>2</sub>R<sup>9</sup> and -S(O)<sub>s</sub>R<sup>9</sup>, and A is optionally further substituted by one substituent selected from -OR<sup>7</sup>, halogen, trifluoromethyl, -CN, -CO<sub>2</sub>R<sup>7</sup> and C<sub>1-6</sub>alkyl optionally substituted by hydroxy;

R<sup>1</sup> is selected from methyl and chloro;

R<sup>2</sup> is selected from -NH-CO-R<sup>11</sup> and -CO-NH-(CH<sub>2</sub>)<sub>t</sub>R<sup>12</sup>;

R<sup>3</sup> is selected from hydrogen, C<sub>1-6</sub>alkyl optionally substituted by up to two OH groups, -(CH<sub>2</sub>)<sub>k</sub>-C<sub>3-7</sub>cycloalkyl, -(CH<sub>2</sub>)<sub>k</sub>phenyl optionally substituted by R<sup>13</sup> and/or R<sup>14</sup> and -(CH<sub>2</sub>)<sub>k</sub>heteroaryl optionally substituted by R<sup>13</sup> and/or R<sup>14</sup>,

R<sup>4</sup> is selected from hydrogen and C<sub>1-6</sub>alkyl, or

R<sup>3</sup> and R<sup>4</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R<sup>15</sup>;

R<sup>5</sup> is selected from C<sub>1-6</sub>alkyl optionally substituted by up to three halogen atoms, C<sub>2-6</sub>alkenyl optionally substituted by phenyl, C<sub>3-7</sub>cycloalkyl, heteroaryl optionally substituted by up to three R<sup>13</sup> and/or R<sup>14</sup> groups, and phenyl optionally substituted by R<sup>13</sup> and/or R<sup>14</sup>;

R<sup>6</sup> is a C<sub>3-6</sub>alkyl group substituted by at least two substituents independently selected from -OR<sup>16</sup>, -NR<sup>16</sup>R<sup>17</sup>, -CO<sub>2</sub>R<sup>16</sup>, -CONR<sup>16</sup>R<sup>17</sup>, -NHCOR<sup>16</sup> and -NHSO<sub>2</sub>R<sup>16</sup>;

R<sup>7</sup> and R<sup>8</sup> are each independently selected from hydrogen and C<sub>1-6</sub>alkyl;

R<sup>9</sup> is selected from hydrogen, -(CH<sub>2</sub>)<sub>u</sub>-C<sub>3-7</sub>cycloalkyl, -(CH<sub>2</sub>)<sub>u</sub>heterocyclyl, -(CH<sub>2</sub>)<sub>u</sub>aryl, and C<sub>1-6</sub>alkyl optionally substituted by up to two substituents independently selected from -OR<sup>18</sup> and -NR<sup>18</sup>R<sup>19</sup>,

R<sup>10</sup> is selected from hydrogen and C<sub>1-6</sub>alkyl, or

R<sup>9</sup> and R<sup>10</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R<sup>15</sup>;

R<sup>11</sup> is selected from hydrogen, C<sub>1-6</sub>alkyl, -(CH<sub>2</sub>)<sub>t</sub>-C<sub>3-7</sub>cycloalkyl, trifluoromethyl, -(CH<sub>2</sub>)<sub>t</sub>heteroaryl optionally substituted by R<sup>20</sup> and/or R<sup>21</sup>, and -(CH<sub>2</sub>)<sub>v</sub>phenyl optionally substituted by R<sup>20</sup> and/or R<sup>21</sup>;

R<sup>12</sup> is selected from hydrogen, C<sub>1-6</sub>alkyl, C<sub>3-7</sub>cycloalkyl, -CONHR<sup>22</sup>, phenyl optionally substituted by R<sup>20</sup> and/or R<sup>21</sup>, and heteroaryl optionally substituted by R<sup>20</sup> and/or R<sup>21</sup>;

R<sup>13</sup> and R<sup>14</sup> are each independently selected from halogen, -CN, trifluoromethyl, nitro, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, -CONR<sup>22</sup>R<sup>23</sup>, -COR<sup>24</sup>, -CO<sub>2</sub>R<sup>24</sup>, and heteroaryl, or

R<sup>13</sup> and R<sup>14</sup> are linked to form a fused 5-membered heterocyclyl ring containing one heteroatom selected from oxygen, sulfur and N-R<sup>15</sup>, or a fused heteroaryl ring;

R<sup>15</sup> is selected from hydrogen and methyl;

R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup> and R<sup>19</sup> are each independently selected from hydrogen and C<sub>1-6</sub>alkyl;

R<sup>20</sup> is selected from C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, -(CH<sub>2</sub>)<sub>t</sub>-C<sub>3-7</sub>cycloalkyl, -CONR<sup>22</sup>R<sup>23</sup>, -NHCOR<sup>23</sup>, halogen, -CN, -(CH<sub>2</sub>)<sub>w</sub>NR<sup>25</sup>R<sup>26</sup>, trifluoromethyl, phenyl optionally substituted by one or more R<sup>21</sup> groups, and heteroaryl optionally substituted by one or more R<sup>21</sup> groups;

R<sup>21</sup> is selected from C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, halogen, trifluoromethyl, and -(CH<sub>2</sub>)<sub>w</sub>NR<sup>25</sup>R<sup>26</sup>;

R<sup>22</sup> and R<sup>23</sup> are each independently selected from hydrogen and C<sub>1-6</sub>alkyl, or

R<sup>22</sup> and R<sup>23</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R<sup>15</sup>, wherein the ring may be substituted by up to two C<sub>1-6</sub>alkyl groups;

R<sup>24</sup> is C<sub>1-6</sub>alkyl;

R<sup>25</sup> is selected from hydrogen, C<sub>1-6</sub>alkyl and -(CH<sub>2</sub>)<sub>t</sub>-C<sub>3-7</sub>cycloalkyl optionally substituted by C<sub>1-6</sub>alkyl,

R<sup>26</sup> is selected from hydrogen and C<sub>1-6</sub>alkyl, or

R<sup>25</sup> and R<sup>26</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R<sup>15</sup>;

R<sup>27</sup> is hydrogen or C<sub>1-6</sub>alkyl;

[[B]] B<sub>1</sub> is selected from a bond, oxygen, NH and S(O)<sub>x</sub>;

X and Y are each independently selected from hydrogen, methyl and halogen;

Z is selected from halogen, C<sub>1-6</sub>alkyl and -OR<sup>27</sup>;

k, m and w are each independently selected from 0, 1, 2 and 3;

n, q, r, s, t and x are each independently selected from 0, 1 and 2; and

u and v are each independently selected from 0 and 1;  
or a pharmaceutically acceptable ~~derivative~~ salt thereof.

2. (original) A compound according to claim 1 wherein A is a fused 5-membered heteroaryl ring containing up to two heteroatoms independently selected from oxygen and nitrogen.

3. (previously presented) A compound according to claim 1 wherein A is substituted by  $-(CH_2)_q$ aryl or  $-(CH_2)_q$ heteroaryl wherein the aryl or heteroaryl is optionally substituted by one or more substituents independently selected from oxo,  $C_{1-6}$ alkyl, halogen,  $-CN$ , trifluoromethyl,  $-OR^9$ ,  $-(CH_2)_tCO_2R^{10}$ ,  $-NR^9R^{10}$ ,  $-(CH_2)_tCONR^9R^{10}$ ,  $-NHCOR^9$ ,  $-SO_2NR^9R^{10}$ ,  $-NHSO_2R^9$  and  $-S(O)_sR^9$ .

4. (previously presented) A compound according to claim 1 wherein  $R^1$  is methyl.

5. (previously presented) A compound according to claim 1 wherein  $R^2$  is  $-CO-NH-(CH_2)_t-R^{12}$ .

6. (previously presented) A compound according to claim 1 wherein X is hydrogen or fluorine.

7. (currently amended) A compound according to claim 1 which is  
N-Cyclopropyl-3-[5-fluoro-3-(4-pyridinyl)-1H-indazol-6-yl]-4-methylbenzamide;  
N-Cyclopropyl-3-[5-fluoro-3-(1-oxido-4-pyridinyl)-1H-indazol-6-yl]-4-methylbenzamide;  
N-Cyclopropyl-3-fluoro-5-[5-fluoro-3-(4-pyridinyl)-1,2-benzisoxazol-6-yl]-4-  
methylbenzamide;  
N-Cyclopropyl-3-fluoro-5-[5-fluoro-3-(1-oxido-4-pyridinyl)-1,2-benzisoxazol-6-yl]-4-  
methylbenzamide;  
N-Ethyl-3-[5-fluoro-3-[6-(methyloxy)-3-pyridinyl]-1H-indazol-6-yl]-4-methylbenzamide;  
3-[3-(6-Chloro-3-pyridinyl)-5-fluoro-1H-indazol-6-yl]-N-ethyl-4-methylbenzamide;

~~substantially as hereinbefore defined with reference to any one of Examples 1 to 6, or a~~  
pharmaceutically acceptable ~~derivative~~ salt thereof.

8. (Currently amended) A compound ~~selected from~~ which is:

*N*-cyclopropyl-3-[5-fluoro-3-(4-pyridinyl)-1*H*-indazol-6-yl]-4-methylbenzamide; ~~or~~ and  
*N*-cyclopropyl-3-fluoro-5-[5-fluoro-3-(4-pyridinyl)-1,2-benzisoxazol-6-yl]-4-  
methylbenzamide;

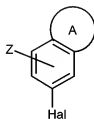
or a pharmaceutically acceptable derivative thereof.

9. (previously presented) A pharmaceutical composition comprising at least one  
compound as claimed in claim 1, or a pharmaceutically acceptable derivative thereof, in  
association with one or more pharmaceutically acceptable excipients, diluents and/or carriers.

10 to 13. (cancelled)

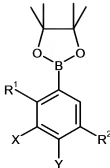
14. (currently amended) A process for preparing a compound of formula (I) as claimed  
in claim 1, or a pharmaceutically acceptable ~~derivative~~ salt thereof, which comprises

(a) reacting a compound of formula (II)

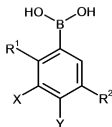


(II)

in which A is defined in claim 1 and Hal is halogen,  
with a compound of formula (IIIA) or (IIIB)



(IIIA)



(IIIB)

in which R<sup>1</sup>, R<sup>2</sup>, X and Y are as defined in claim 1,  
in the presence of a catalyst, or

- (b) final stage modification of one compound of formula (I) as defined in claim 1 to give another compound of formula (I) as defined in claim 1.

15. (previously presented) A compound according to claim 3 wherein A is substituted by  $-(CH_2)_q$ heteroaryl wherein the heteroaryl is optionally substituted by one or more substituents independently selected from oxo, C<sub>1-6</sub>alkyl, halogen, -CN, trifluoromethyl, -OR<sup>9</sup>,  $-(CH_2)_tCO_2R^{10}$ , -NR<sup>9</sup>R<sup>10</sup>,  $-(CH_2)_tCONR^9R^{10}$ , -NHCOR<sup>9</sup>, -SO<sub>2</sub>NR<sup>9</sup>R<sup>10</sup>, -NHSO<sub>2</sub>R<sup>9</sup> and -S(O)<sub>s</sub>R<sup>9</sup>.

16. (previously presented) A compound according to claim 15 wherein R<sup>1</sup> is methyl.

17. (previously presented) A compound according to claim 15 wherein R<sup>2</sup> is -CO-NH-(CH<sub>2</sub>)<sub>t</sub>-R<sup>12</sup>.

18. (previously presented) A compound according to claim 15 wherein X is hydrogen or fluorine.

19 (Currently amended). A compound according to Claim 15 wherein the 5-membered ring fused to the phenyl ring is an optionally substituted indazole.

20. (previously presented) A compound according to Claim 15 wherein the heteroaryl is a 5- or 6-membered heteroaryl ring containing up to two heteroatoms independently selected from oxygen and nitrogen.

21. (previously presented) A compound according to Claim 20 wherein the heteroaryl ring is a pyridyl.

22. (previously presented) A compound according to Claim 21 wherein q is 0.

23. (previously presented) A compound according to Claim 1 wherein Z is a halogen.

24 (new). A compound according to Claim 1 wherein the 5-membered ring A fused to the phenyl ring is an optionally substituted isoxazolyl, indazole, pyrazolyl or pyrrolyl.